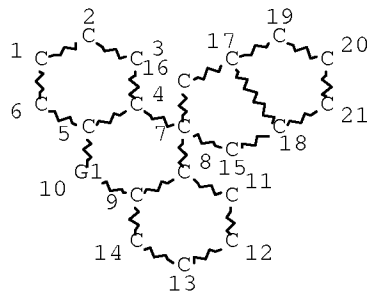


10/556,717

=> d que 137

L3 STR



REP G1=(0-3) CH2

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DEFAULT ECLEVEL IS LIMITED

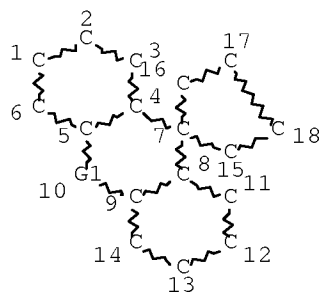
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STEREO ATTRIBUTES: NONE

L5 STR



REP G1=(0-3) CH2

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

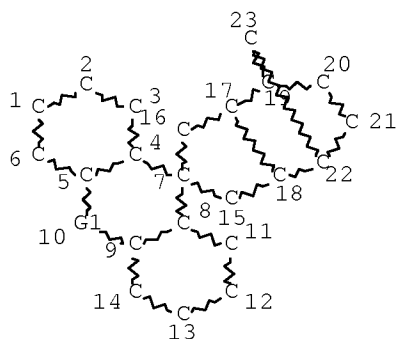
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RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 18

STEREO ATTRIBUTES: NONE

L9 STR



REP G1=(0-3) CH2

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DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

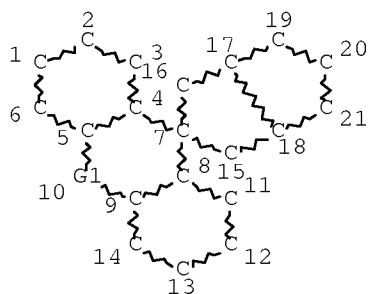
NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

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L24	54	SEA FILE=REGISTRY	SUB=L11 SSS FUL	L3
L26	46	SEA FILE=REGISTRY	ABB=ON PLU=ON	L24 NOT FULLER?
L27	35	SEA FILE=REGISTRY	ABB=ON PLU=ON	L26 NOT FLUORANTHENE?
L28	34	SEA FILE=REGISTRY	ABB=ON PLU=ON	L27 NOT 1-100/SI
L29	30	SEA FILE=REGISTRY	ABB=ON PLU=ON	L28 NOT DISPIRO?
L32	29	SEA FILE=REGISTRY	ABB=ON PLU=ON	L29 NOT MORPHOLINYL?
L33	28	SEA FILE=REGISTRY	ABB=ON PLU=ON	L32 NOT MORPHOLINE?
L34	15	SEA FILE=REGISTRY	ABB=ON PLU=ON	L33 AND L14
L36	14	SEA FILE=HCAPLUS	ABB=ON PLU=ON	L34
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=> d que 138

L3 STR



REP G1=(0-3) CH2

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

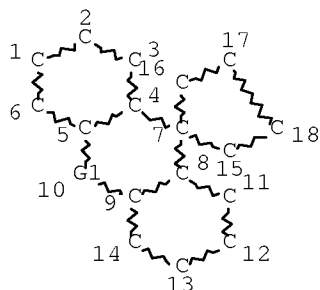
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RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 21

STEREO ATTRIBUTES: NONE

L5 STR



REP G1=(0-3) CH2

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

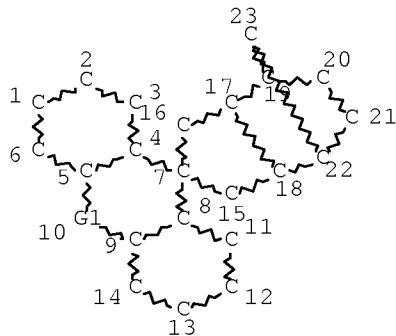
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RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 18

STEREO ATTRIBUTES: NONE

L9 STR



REP G1=(0-3) CH2

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

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10/556,717

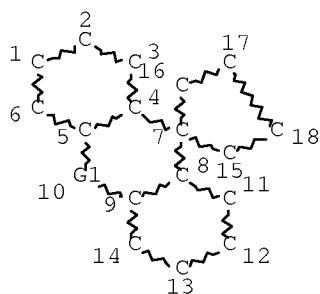
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L24      54 SEA FILE=REGISTRY SUB=L11 SSS FUL L3
L26      46 SEA FILE=REGISTRY ABB=ON  PLU=ON  L24 NOT FULLER?
L27      35 SEA FILE=REGISTRY ABB=ON  PLU=ON  L26 NOT FLUORANTHENE?
L28      34 SEA FILE=REGISTRY ABB=ON  PLU=ON  L27 NOT 1-100/SI
L29      30 SEA FILE=REGISTRY ABB=ON  PLU=ON  L28 NOT DISPIRO?
L32      29 SEA FILE=REGISTRY ABB=ON  PLU=ON  L29 NOT MORPHOLINYL?
L33      28 SEA FILE=REGISTRY ABB=ON  PLU=ON  L32 NOT MORPHOLINE?
L34      15 SEA FILE=REGISTRY ABB=ON  PLU=ON  L33 AND L14
L36      14 SEA FILE=HCAPLUS ABB=ON  PLU=ON  L34
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L38      0  SEA FILE=MARPAT ABB=ON  PLU=ON  L20 NOT L37

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=> d que 116

L5 STR



REP G1=(0-3) CH2

NODE ATTRIBUTES:

DEFAULT MLEVEL IS ATOM

DEFAULT ECLEVEL IS LIMITED

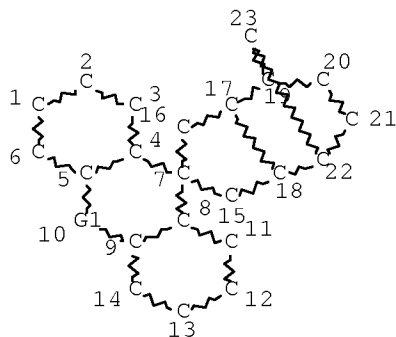
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RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 18

STEREO ATTRIBUTES: NONE

L9 STR



REP G1=(0-3) CH2

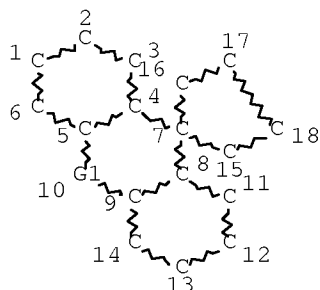
10/556,717

NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
RING(S) ARE ISOLATED OR EMBEDDED
NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE
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L16 0 SEA FILE=CAOLD ABB=ON PLU=ON L14

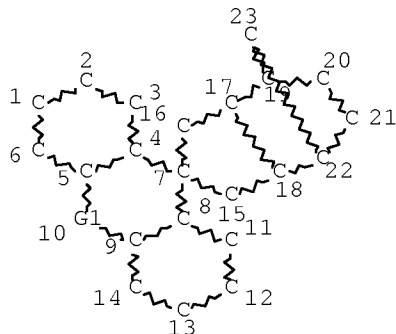
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L5 STR



REP G1=(0-3) CH2
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DEFAULT MLEVEL IS ATOM
DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:
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NUMBER OF NODES IS 18

STEREO ATTRIBUTES: NONE
L9 STR



REP G1=(0-3) CH2
NODE ATTRIBUTES:
DEFAULT MLEVEL IS ATOM

10/556,717

DEFAULT ECLEVEL IS LIMITED

GRAPH ATTRIBUTES:

RING(S) ARE ISOLATED OR EMBEDDED

NUMBER OF NODES IS 23

STEREO ATTRIBUTES: NONE

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L14 15 SEA FILE=REGISTRY SUB=L11 SSS FUL L9

L18 0 SEA FILE=BEILSTEIN ABB=ON PLU=ON L14

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L37 ANSWER 1 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:583235 HCAPLUS Full-text
 DOCUMENT NUMBER: 148:540349
 TITLE: Purification of cycloolefin compounds
 INVENTOR(S): Kaizu, Michitaka; Masutani, Masataka; Oda,
 Hirokazu; Nishimura, Toshiro
 PATENT ASSIGNEE(S): Jsr Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 14pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
-----	----	-----	-----	-----
JP 2008110930	A	20080515	JP 2006-294259	20061030
PRIORITY APPLN. INFO.:			JP 2006-294259	20061030

ED Entered STN: 15 May 2008

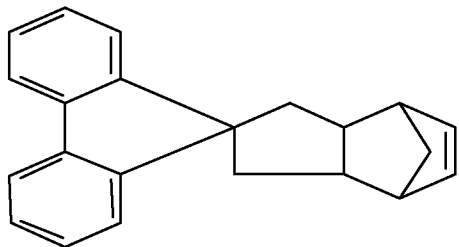
AB Cycloolefin compds. are purified by (1) contacting organic solvent A solution of cycloolefin compds. with aqueous medium to remove water-soluble components from A to the aqueous medium, (2) removing at least a part of A, and (3) adding a mixture of A and poor solvent B for the cycloolefin compds. to the concentrated solution for precipitation of the products. Thus, a reaction product containing spiro[fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene] (I) was washed with H₂O and the toluene layer was concentrated to half and added dropwise to a mixture of toluene and MeOH. The mixture was further stirred at 0° for 1 h to give 67.2% I with purity 99.24%.

IT 908004-25-1P

(purification of cycloolefin compds. by removing water-soluble components from organic solvent solution upon washing with aqueous medium, concentration, and precipitation with poor solvent)

RN 908004-25-1 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro- (CA INDEX NAME)



CC 45-4 (Industrial Organic Chemicals, Leather, Fats, and Waxes)

IT 908004-25-1P

(purification of cycloolefin compds. by removing water-soluble components from organic solvent solution upon washing with aqueous medium, concentration, and precipitation with poor solvent)

L37 ANSWER 2 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:468745 HCAPLUS Full-text

DOCUMENT NUMBER: 148:450087

TITLE: Manufacture of spiro hydrocarbons with low cost of organic solvents

INVENTOR(S): Kaizu, Michitaka; Masutani, Masataka; Nishimura, Toshiro

PATENT ASSIGNEE(S): JSR Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 20pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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JP 2008088091	A	20080417	JP 2006-269717	20060929
PRIORITY APPLN. INFO.:			JP 2006-269717	20060929

ED Entered STN: 17 Apr 2008

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Spiro hydrocarbons I [R1-R8 = H, polar group, C1-30 hydrocarbyl which may have C1-10 hydrocarbylene or linking group containing O, N, S, or Si; R10, R11 = H, polar group, C1-30 hydrocarbyl including (un)substituted unsatd. structure which may have linking group containing O, N, S, or Si; R12-R17 = H, C1-10 hydrocarbylene; A = single bond, O, S, SO, SO2, CO, (CH2)n, NR18, Si(R19)2; R18, R19 = C1-30 hydrocarbyl which may have halogen atom; a-c, n = 0-2] or II (R1-R15, A, a-c = same as above), useful for polymeric optical materials, are manufactured by feeding cyclic hydrocarbons III (R1-R8, A, a-c = same as above) and alkali metal hydroxides into organic solvents having b.p. 100-200°, and mixing with solns. comprising the organic solvents and XCR12R13CR10R16CR11R17CR14R15Y [IV; R10-R17 = same as above; X, Y = halo, R20SO3, (R21O)R22P(O)O; R21, R22 = (halo-substituted) C1-30 hydrocarbyl] or XCR12R13CR10:CR11CR14R15Y (V; R10-R15, X, Y = same as above) under reflux conditions. Preferably, ≥30% of the organic solvents are recovered from reaction mixts. by distillation, and recycled to manufacture of IV or V, or to manufacture of I or II. Thus, a solution comprising PhMe 446, fluorene 65.3, KOH 97.9 kg was heated, gradually mixed with a solution comprising 549 kg PhMe and 182 kg 5,6-di(p- toluenesulfonyloxymethyl)bicyclo[2.2.1]hept-2-ene (VI) under reflux condition, and washed with water 4 times to give a PhMe phase containing spiro(fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene) in 76.2% yield. Then, 500 kg PhMe was recovered from the PhMe phase. VI was manufactured by using the recovered PhMe PhMe in a yield comparable to that in manufacture by using virgin PhMe.

IT 908004-25-1P

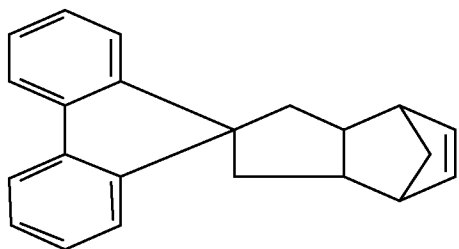
(manufacture of spiro hydrocarbons by cyclization of hydrocarbons with

10/556,717

active methylene compds. and in-process recycling of organic solvents)

RN 908004-25-1 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro- (CA INDEX NAME)



CC 35-2 (Chemistry of Synthetic High Polymers)

Section cross-reference(s): 25, 73

IT 908004-25-1P

(manufacture of spiro hydrocarbons by cyclization of hydrocarbons with active methylene compds. and in-process recycling of organic solvents)

L37 ANSWER 3 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:416735 HCAPLUS [Full-text](#)

DOCUMENT NUMBER: 148:404056

TITLE: Norbornene-based polymers with negative retardation in the thickness direction and films useful for polarizing plate and liquid crystal display device made from the polymers

INVENTOR(S): Nozoe, Yutaka; Watanabe, Saisuke

PATENT ASSIGNEE(S): Fujifilm Corporation, Japan

SOURCE: U.S. Pat. Appl. Publ., 18pp.

CODEN: USXXCO

DOCUMENT TYPE: Patent

LANGUAGE: English

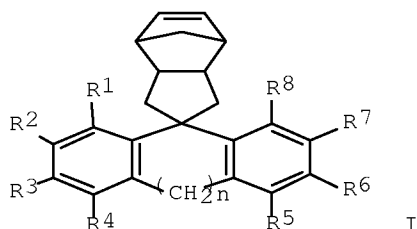
FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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US 20080081890	A1	20080403	US 2007-862322	20070927
US 7390864	B2	20080624		
JP 2008081655	A	20080410	JP 2006-265002	20060928
KR 2008029834	A	20080403	KR 2007-97241	20070927
PRIORITY APPLN. INFO.:			JP 2006-265002	A 20060928

ED Entered STN: 03 Apr 2008

GI



AB The polymers are formed by addition polymerization of a composition comprising at least one norbornene derivative represented by I (R1, R2, R3, R4, R5, R6, R7, R8 = H or a substituent; n = 0 to 2). Thus, heating dicyclopentadiene with allyl acetate in the presence of small amount of hydroquinone at 180° for 9 h gave a product which was polymerized to give a polymer having the claimed properties.

IT 1015787-82-2P 1015787-83-3P 1015787-85-5P
(norbornene-based polymers with neg. retardation in thickness direction and films useful for polarizing plate and liquid crystal display device made from the polymers)

RN 1015787-82-2 HCAPLUS

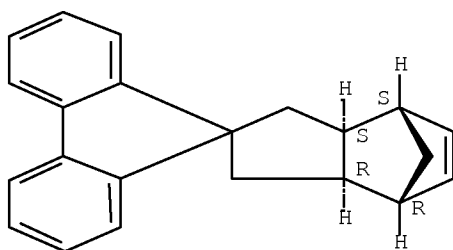
CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'R,7'S,7'aS)-rel-, homopolymer (CA INDEX NAME)

CM 1

CRN 791813-27-9

CMF C22 H20

Relative stereochemistry.



RN 1015787-83-3 HCAPLUS

CN Bicyclo[2.2.1]hept-5-ene-2-methanol, 2-acetate, polymer with rel-(3'aR,4'R,7'S,7'aS)-1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] (CA INDEX NAME)

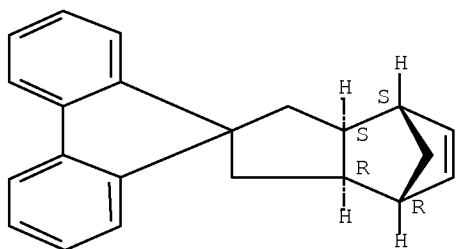
CM 1

CRN 791813-27-9

CMF C22 H20

Relative stereochemistry.

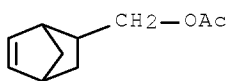
10/556,717



CM 2

CRN 10471-24-6

CMF C10 H14 O2



RN 1015787-85-5 HCAPLUS

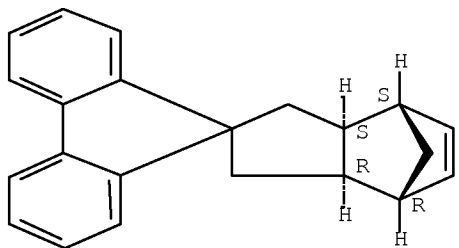
CN Pentanoic acid, bicyclo[2.2.1]hept-5-en-2-ylmethyl ester, polymer with
rel-(3'aR,4'R,7'S,7'aS)-1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-
9,2'-[4,7]methano[2H]indene] (CA INDEX NAME)

CM 1

CRN 791813-27-9

CMF C22 H20

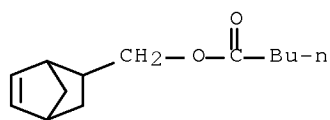
Relative stereochemistry.



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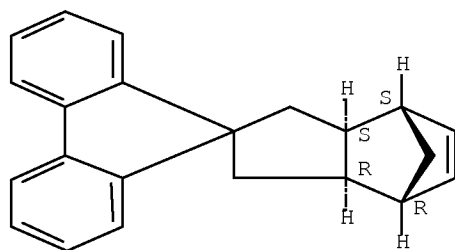
CRN 118361-19-6

CMF C13 H20 O2



IT 791813-27-9P
 (norbornene-based polymers with neg. retardation in thickness
 direction and films useful for polarizing plate and liquid crystal
 display device made from the polymers)
 RN 791813-27-9 HCAPLUS
 CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-
 hexahydro-, (3'aR,4'R,7'S,7'aS)-rel- (CA INDEX NAME)

Relative stereochemistry.



INCL -526
 CC 37-3 (Plastics Manufacture and Processing)
 Section cross-reference(s): 38, 74
 IT 26935-85-3P 1015787-82-2P 1015787-83-3P
 1015787-85-5P
 (norbornene-based polymers with neg. retardation in thickness
 direction and films useful for polarizing plate and liquid crystal
 display device made from the polymers)
 IT 6143-29-9P 70096-09-2P 118361-19-6P 791813-27-9P
 (norbornene-based polymers with neg. retardation in thickness
 direction and films useful for polarizing plate and liquid crystal
 display device made from the polymers)
 REFERENCE COUNT: 5 THERE ARE 5 CITED REFERENCES AVAILABLE FOR
 THIS RECORD. ALL CITATIONS AVAILABLE IN THE
 RE FORMAT

L37 ANSWER 4 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2008:278915 HCAPLUS Full-text
 DOCUMENT NUMBER: 148:309101
 TITLE: Cycloolefin copolymers with uniform distribution
 of structural units, and their manufacture method
 and uses as optical materials
 INVENTOR(S): Yoshida, Shuichi; Mutsuga, Yasuaki; Kawashima,
 Naoyuki; Miyaki, Nobuyuki; Kajiwara, Ichiro
 PATENT ASSIGNEE(S): Jsr Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 29pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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JP 2008050495	A	20080306	JP 2006-229592	20060825
PRIORITY APPLN. INFO.:			JP 2006-229592	20060825

ED Entered STN: 06 Mar 2008

AB Title method involves copolymn. of (A) cycloolefins having norbornene skeletons and aromatic ring structures with (B) cycloolefins having norbornene skeletons and no aromatic ring structures, wherein the polymerization is started using 5-90% of total monomers and polymerization catalysts and the residual monomers are added to the system during the polymerization. The copolymers with good transparency, heat resistance, solubility in organic solvents, strength, and processability, show (maldistribution of the aromatic ring structures) = $| (R100000)/(U100000) - (R1000)/(U1000) | \leq 1$ (R100000 and R1000 are intensities at mol. weight 10,000 and 1000 in differential mol. weight distribution curves obtained from refractive index measurements, resp.; U100000 and U1000 are intensities at mol. weight 100,000 and 1000 in differential mol. weight distribution curves obtained from UV absorption measurements at 254 nm, resp.). The uses include optical parts, (stretched) optical films showing peculiar birefringence and wavelength dependence, polarizing plates with the stretched films, and liquid crystal displays equipped with the polarizing plates. Thus, copolymg. spiro[fluorene-9,8'-tricyclo[4.3.0.12,5][3]decene] 22, 8-methoxycarbonyl-8-methyltetracyclo[4.4.0.12,5.17,10]-3-dodecene (I) 59, and bicyclo[2.2.1]hept-2-ene 4 g in the presence of Et₃Al and WCl₆, dropwise adding 15 g I after temperature increase rate of the polymerization solution reached 1°/min, further polymerizing, and hydrogenating gave a hydrogenated copolymer (maldistribution of the aromatic ring structures 0.74, logarithmic viscosity 0.72, T_g 165°), which was cast on a glass container, released from the glass surfaces, and uniaxially stretched to give a stretched film showing maximum stress 64 kg/cm² at 2-fold stretching, haze 0.4, retardation 210, 235, and 244 nm at 450, 550, and 650 nm, resp. A liquid crystal TV with the stretched film showed good contrast.

IT 1009643-07-SDP, hydrogenated
(cycloolefin copolymers with suppressed uniform distribution of structural units and their optical films for polarizing plates of liquid crystal displays)

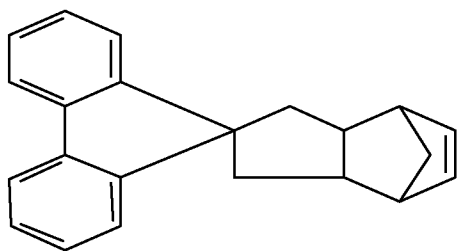
RN 1009643-07-5 HCAPLUS

CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-octahydro-2-methyl-, methylester, polymer with 1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] and bicyclo[2.2.1]hept-2-ene (CA INDEX NAME)

CM 1

CRN 908004-25-1

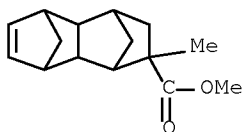
CMF C22 H20



CM 2

CRN 58732-15-3

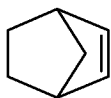
CMF C15 H20 O2



CM 3

CRN 498-66-8

CMF C7 H10



CC 37-3 (Plastics Manufacture and Processing)

Section cross-reference(s): 38, 73, 74

IT 1009643-07-5DP, hydrogenated

(cycloolefin copolymers with suppressed uniform distribution of structural units and their optical films for polarizing plates of liquid crystal displays)

L37 ANSWER 5 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2008:249623 HCAPLUS [Full-text](#)

DOCUMENT NUMBER: 148:286392

TITLE: Cyclic olefin addition copolymers, their manufacture, and their optical members and films
 INVENTOR(S): Sakagami, Toshinori; Hirano, Akira; Maruyama, Yoichiro

PATENT ASSIGNEE(S): Jsr Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 43pp.

DOCUMENT TYPE: CODEN: JKXXAF
 LANGUAGE: Patent
 FAMILY ACC. NUM. COUNT: 1 Japanese
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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JP 2008045069	A	20080228	JP 2006-223512	20060818
PRIORITY APPLN. INFO.:			JP 2006-223512	20060818

ED Entered STN: 28 Feb 2008
 GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

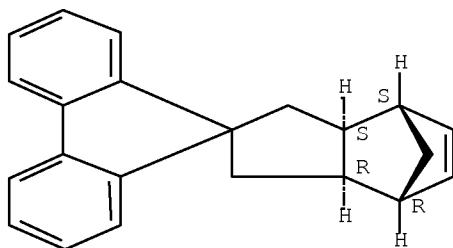
AB The copolymers are manufactured by addition copolymn. of I (A1-A4 = H, halo, C1-20 alkyl, aryl, cycloalkyl, halogenated hydrocarbon, hydrolyzable silyl group, (CH₂)_qX; A1 and A2 or A3 and A4 may link together to form an alkylidene group; A1 and A2, A3 and A4 or A2 and A3 may link together to form C or heterocyclic rings; p = 0, 1; q = 0-5; X = CO₂A₅, OCOA₆, OA₇; A₅, A₆, A₇ = C1-10 hydrocarbon group, halogenated hydrocarbon, oxetanyl-containing substituent) and II [m, n = 0-2; B1-B8 = H, halo, (linkage group-containing) C1-30 hydrocarbon group, polar group; s, t, u = 0-3] in the presence of polymerization catalysts (A) Pd organic acid salts or β-diketonate compds., (B) phosphine compds. PR₁2R₂ (R₁ = cyclopentyl, cyclohexyl, isopropyl; R₂ = C3-10 hydrocarbon group), and (D) ionic B compds., or (C) divalent Pd phosphine complexes Pd(PR₁2R₂)_nX₂ (R₁, R₂ = same as above; X = organic acid anion or β-diketonate anion; n = 1, 2), and D. Thus, polymerization of 5-butylobicyclo[2.2.1]hepta-2-ene, 5-decylobicyclo[2.2.1]hepta-2-ene, and spiro[fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene] in the presence of (tricyclopentylphosphine)palladium diacetate and triphenylcarbenium tetrakis(pentafluorophenyl) borate gave a copolymer, which was mixed with additives and vacuum-pressed to give a film with T_g 232° and low retardation and water absorption.

IT 791813-27-9P, exo-Spiro[fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene]
 (manufacture of cyclic olefin addition copolymers for optical films with low retardation)

RN 791813-27-9 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'R,7'S,7'aS)-rel- (CA INDEX NAME)

Relative stereochemistry.



IT 1008099-98-6P, 5-Butylbicyclo[2.2.1]hepta-2-ene-5-decylbicyclo[2.2.1]hepta-2-ene-exo-spiro[fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene] copolymer 1008100-00-2P, Bicyclo[2.2.1]hepta-2-ene-5-butylbicyclo[2.2.1]hepta-2-ene-exo-spiro(fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene) copolymer (manufacture of cyclic olefin addition copolymers for optical films with low retardation)

RN 1008099-98-6 HCAPLUS

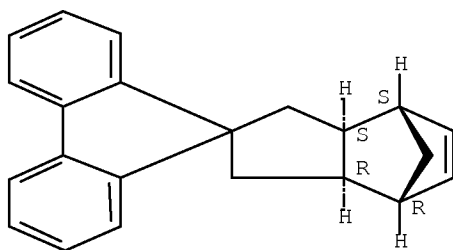
CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'R,7'S,7'aS)-rel-, polymer with 5-butylbicyclo[2.2.1]hept-2-ene and 5-decylbicyclo[2.2.1]hept-2-ene (CA INDEX NAME)

CM 1

CRN 791813-27-9

CMF C22 H20

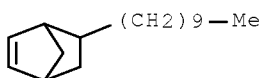
Relative stereochemistry.



CM 2

CRN 22094-85-5

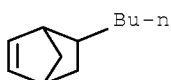
CMF C17 H30



CM 3

CRN 22094-81-1

CMF C11 H18



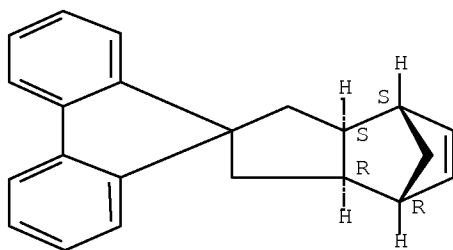
RN 1008100-00-2 HCAPLUS
 CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'R,7'S,7'aS)-rel-, polymer with bicyclo[2.2.1]hept-2-ene and 5-butylbicyclo[2.2.1]hept-2-ene (CA INDEX NAME)

CM 1

CRN 791813-27-9

CMF C22 H20

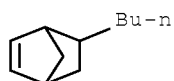
Relative stereochemistry.



CM 2

CRN 22094-81-1

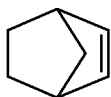
CMF C11 H18



CM 3

CRN 498-66-8

CMF C7 H10



CC 38-3 (Plastics Fabrication and Uses)
 Section cross-reference(s): 25, 35, 73
 IT 791813-27-9F, exo-Spiro[fluorene-9,8'-

tricyclo[4.3.0.12.5][3]decene]

(manufacture of cyclic olefin addition copolymers for optical films with low retardation)

IT 1008099-98-6P, 5-Butylbicyclo[2.2.1]hepta-2-ene-5-decylbicyclo[2.2.1]hepta-2-ene-exo-spiro[fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene] copolymer 1008100-00-2P, Bicyclo[2.2.1]hepta-2-ene-5-butylbicyclo[2.2.1]hepta-2-ene-exo-spiro(fluorene-9,8'-tricyclo[4.3.0.12.5][3]decene) copolymer (manufacture of cyclic olefin addition copolymers for optical films with low retardation)

L37 ANSWER 6 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:1237371 HCAPLUS Full-text

DOCUMENT NUMBER: 147:502739

TITLE: Preparation of polycyclic hydrocarbon compounds

INVENTOR(S): Hirano, Akira; Katsuta, Kohei; Miyaki, Nobuyuki; Ebata, Satoshi

PATENT ASSIGNEE(S): Jsr Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 25pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

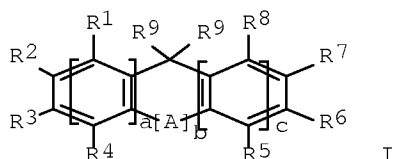
PATENT INFORMATION:

PATENT NO. -----	KIND ----	DATE -----	APPLICATION NO. -----	DATE -----
JP 2007284401	A	20071101	JP 2006-115582	20060419
PRIORITY APPLN. INFO.:			JP 2006-115582	20060419

OTHER SOURCE(S): MARPAT 147:502739

ED Entered STN: 01 Nov 2007

GI



AB Title compds. (I) [R1-R8 = H, halo, (un)substituted C1-30 hydrocarbyl optionally having C1-10 divalent linking group or linking group containing O, N, S, or Si, polar group; R9 = H, halo, C1-30 hydrocarbyl optionally having linking group containing O, N, S, or Si, polar group; 2 R9 groups may be bonded together to form CH2CHR10CHR11CH2, CH2CR10:CR11CH2; R10, R11 = any group given for R9; R10 and R11 may be bonded together to form (hetero)cyclic ring; a, b, c = 0-2; A = direct bond, O, S, SO, SO2, CO, NR18, SiR192 [R18, R19 = C1-30 (halo)hydrocarbyl], (CH2)m (n = 0-2)], useful as monomers or monomer precursors for cycloolefin polymers, are prepared by reacting I (R1-R9, A, a, b, c = same as above; R9 = H) with R9X [R9 = same as above; X = halo, OSO2R20 [R20 = C1-30 (halo)hydrocarbyl], OP(O)R22(OR21) [R21, R22 = C1-30 (halo)hydrocarbyl]], XCH2CHR10CHR11CH2Y [R10, R11 = same as above; X, Y = any group given for X; X and Y may be bonded together to form OP(O)R22O], or

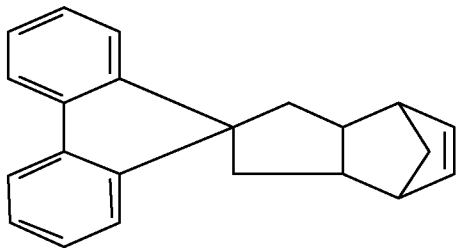
XCH2CR10:CR11CH2Y in hydrocarbon solvents in the presence of alkali metal hydroxides. Thus, KOH was gradually added to toluene solution of 5,6-di(p-toluenesulfonyloxymethyl)bicyclo[2. 2.1]hept-2-ene and fluorene under reflux. Yield of spiro[fluorene-9,8'-tricyclo[4.3.0.12,5][3]decene] 5 h after addition of KOH was 58%.

IT 908004-25-1F

(preparation of (spiro structure-containing) polycyclic hydrocarbon compds. as monomers (precursors) for cycloolefin polymers)

RN 908004-25-1 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro- (CA INDEX NAME)



CC 35-2 (Chemistry of Synthetic High Polymers)

IT 908004-25-1F

(preparation of (spiro structure-containing) polycyclic hydrocarbon compds. as monomers (precursors) for cycloolefin polymers)

L37 ANSWER 7 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2007:760479 HCAPLUS [Full-text](#)

DOCUMENT NUMBER: 147:167069

TITLE: Ring-opening polymerization polymer of cycloolefin, optical parts and films therefrom

INVENTOR(S): Kawashima, Naoyuki; Miyaki, Nobuyuki; Miyamoto, Yoshikazu; Kajiwara, Ichiro

PATENT ASSIGNEE(S): JSR Corporation, Japan

SOURCE: PCT Int. Appl., 58pp.

CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

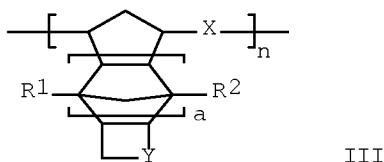
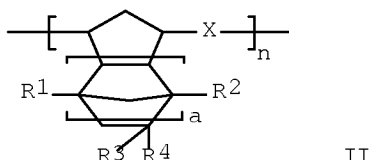
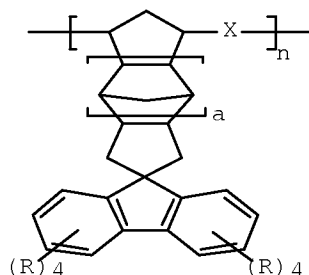
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007077771	A1	20070712	WO 2006-JP325661	20061222
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, GT, HN, HR, HU, ID, IL, IN, IS, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM,			

10/556,717

ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM
 JP 2007204731 A 20070816 JP 2006-254919 20060920
 PRIORITY APPLN. INFO.: JP 2006-1344 A 20060106
 JP 2006-254919 A 20060920

ED Entered STN: 13 Jul 2007
 GI



AB The polymer, being good in transparency, heat resistance, solubility in organic solvents, strength, and processability, comprises ≥ 2 kinds of structural units selected from structural units represented by formula I, II, and III, wherein $a = 0$ or 1 ; $R, R1-R4 = H$, halogen, $C1-30$ hydrocarbon group containing O, S, N , or Si , and polar group; $X, Y = -CH=CH-$, or $-CH_2CH_2-$. Thus, 100 g 5-norbornene-2,3-dicarboxylic anhydride was reduced by 39.0 g lithium aluminum hydride in THF to give 83 g 5,6-bis(hydroxymethyl)bicyclo[2.2.1]hept-2-ene, 36 g of which was reacted with 99 g p-toluenesulfonyl chloride to give 66 g 5,6-di(4-toluenesulfonyloxymethyl)bicyclo[2.2.1]hept-2-ene (A). 36 g Fluorene was added with 270 mL 1.6 M Bu lithium in hexane at -50° , added with 50 g A at -50° , stirred at room temperature for 15 h to give 11 g spiro[fluorene-9, 8'-tricyclo[4.3.0.12.5][3]decene], 30 g of which was polymerized with 66 g Me 4-methyltetracyclo[6.2.1.13,6.02,7]dodec-9-ene-4-carboxylate and 4 g 2-norbornylene in the presence of triethylaluminum and methanol-modified WCl_6 at 80° for 1 h to give a polymer, which was hydrogenated in the presence of a catalyst $RuHCl(CO)[P(C_6H_5)_3]_3$ at $160-165^\circ$ to give a title polymer.

IT 943791-16-0 DEF, hydrogenated
 (ring-opening polymerization polymer of cycloolefin, optical parts and films therefrom)

RN 943791-16-0 HCAPLUS

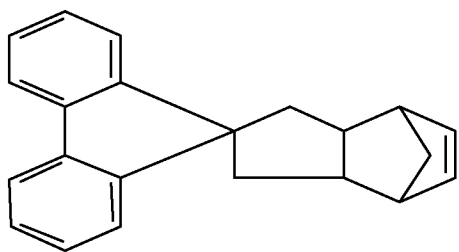
CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-octahydro-, methyl ester, polymer with bicyclo[2.2.1]hept-2-ene and 1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] (CA INDEX NAME)

CM 1

CRN 908004-25-1

CMF C22 H20

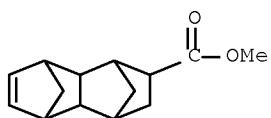
10/556,717



CM 2

CRN 41596-02-5

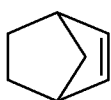
CMF C14 H18 O2



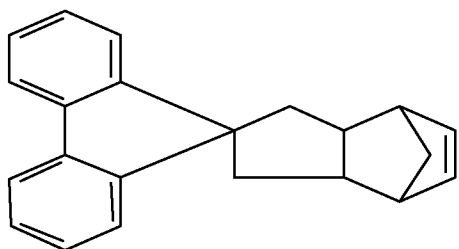
CM 3

CRN 498-66-8

CMF C7 H10



IT 908004-25-1P
 (ring-opening polymerization polymer of cycloolefin, optical parts and
 films therefrom)
 RN 908004-25-1 HCAPLUS
 CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-
 hexahydro- (CA INDEX NAME)



CC 37-3 (Plastics Manufacture and Processing)
 Section cross-reference(s): 38, 73
 IT 943791-16-0DP, hydrogenated
 (ring-opening polymerization polymer of cycloolefin, optical parts and films therefrom)
 IT 85-39-2P, 5,6-Bis(hydroxymethyl)bicyclo[2.2.1]hept-2-ene
 95442-44-7P, 5,6-Di(4-toluenesulfonyloxymethyl)bicyclo[2.2.1]hept-2-ene 908004-25-1P
 (ring-opening polymerization polymer of cycloolefin, optical parts and films therefrom)
 REFERENCE COUNT: 7 THERE ARE 7 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 8 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2007:435619 HCAPLUS [Full-text](#)
 DOCUMENT NUMBER: 146:431510
 TITLE: Process for producing phase difference film, phase difference film, and use thereof
 INVENTOR(S): Mitsuboshi, Tomohiro; Sekiguchi, Masayuki; Ushino, Takuhiro
 PATENT ASSIGNEE(S): JSR Corporation, Japan
 SOURCE: PCT Int. Appl., 72pp.
 CODEN: PIXXD2
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2007043573	A1	20070419	WO 2006-JP320306	20061011
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HN, HR, HU, ID, IL, IN, IS, KE, KG, KM, KN, KP, KR, KZ, LA, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MY, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RS, RU, SC, SD, SE, SG, SK, SL, SM, SV, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, ZA, ZM, ZW RW: AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM				
JP 2007108529	A	20070426	JP 2005-300710	20051014
KR 2008059644	A	20080630	KR 2008-711395	20080513
PRIORITY APPLN. INFO.:			JP 2005-300710	A 20051014

ED Entered STN: 20 Apr 2007

AB The process comprises stretching a norbornene resin film in a film roll widthwise direction by a factor of 1.5-5 and shrinking the film in the longitudinal direction to produce a phase difference film of which the optical axis is in the widthwise direction of the film roll. The phase difference film has excellent transparency, dimensional stability, and view angle properties, has low gas permeability, has an optical axis in the cross direction, and has no significant variation in in-plane phase difference. There is also provided a process for producing a polarizing plate that can produce a high-performance polarizing plate in a simple manner with high production efficiency.

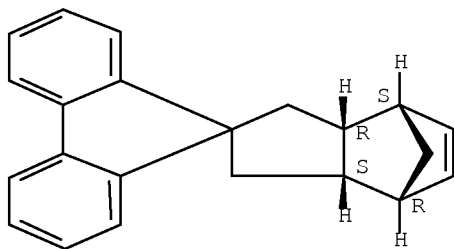
IT 791813-28-0F

(preparation of norbornene derivs. for transparent phase difference films)

RN 791813-28-0 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'S,7'R,7'aS)-rel- (CA INDEX NAME)

Relative stereochemistry.



IT 934389-76-1DF, hydrogenated

(transparent phase difference films for polarizing plates and liquid crystal displays)

RN 934389-76-1 HCAPLUS

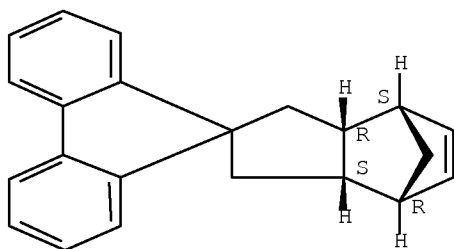
CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-octahydro-2-methyl-, methyl ester, polymer with rel-(3'aR,4'S,7'R,7'aS)-1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] and 1,2,3,4,4a,5,8,8a-octahydro-1,4:5,8-dimethanonaphthalene-2-carboxylic acid (CA INDEX NAME)

CM 1

CRN 791813-28-0

CMF C22 H20

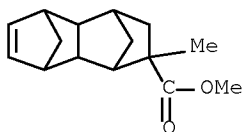
Relative stereochemistry.



CM 2

CRN 58732-15-3

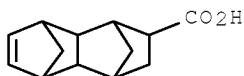
CMF C15 H20 O2



CM 3

CRN 46382-54-1

CMF C13 H16 O2



CC 74-13 (Radiation Chemistry, Photochemistry, and Photographic and Other Reprographic Processes)
 Section cross-reference(s): 38

IT 33493-88-8P 791813-28-0P
 (preparation of norbornene derivs. for transparent phase difference films)

IT 123303-71-9DP, hydrogenated 216987-03-0DP, hydrogenated
 934389-76-1DP, hydrogenated
 (transparent phase difference films for polarizing plates and liquid crystal displays)

REFERENCE COUNT: 4 THERE ARE 4 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 9 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2006:1250985 HCAPLUS Full-text
 DOCUMENT NUMBER: 146:28355

TITLE: Metal hydride complexes for use in hydrogenation of ring-opening polymerization polymer of cycloolefins with freedom from gel complication

INVENTOR(S): Katsuda, Kohei; Fukuhara, Seiji; Maruyama, Yoichiro; Okaniwa, Motoki; Shibata, Hiraku; Suwa, Yoshimi; Kajiwar, Ichiro

PATENT ASSIGNEE(S): JSR Corporation, Japan

SOURCE: PCT Int. Appl., 87pp.
CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2006126612	A1	20061130	WO 2006-JP310398	20060524
W:	AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KM, KN, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, LY, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NG, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SM, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW			
RW:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG, BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM			
JP 2007001967	A	20070111	JP 2005-276253	20050922
JP 2007106932	A	20070426	JP 2005-300459	20051014
EP 1884519	A1	20080206	EP 2006-746827	20060524
R:	AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IS, IT, LI, LT, LU, LV, MC, NL, PL, PT, RO, SE, SI, SK, TR			
CN 101180303	A	20080514	CN 2006-80017733	20071122
KR 2008015026	A	20080215	KR 2007-730082	20071224
PRIORITY APPLN. INFO.:			JP 2005-152163	A 20050525
			JP 2005-276253	A 20050922
			JP 2005-300459	A 20051014
			WO 2006-JP310398	W 20060524

OTHER SOURCE(S): MARPAT 146:28355

ED Entered STN: 30 Nov 2006

AB A metal hydride complex which is a hydride complex of a metal selected from the group consisting of ruthenium, rhodium, osmium, and iridium and has one or more aromatic carboxylic acid residues is provided. It is a novel metal hydride complex which dissolves in a high concentration in organic solvents having low polarity, such as hydrocarbon solvents, and which has high catalytic activity in the hydrogenation of carbon-carbon double bonds. The metal hydride complex is suitable for use as a catalyst especially in the hydrogenation of a ring-opening polymerization polymer of a cycloolefin.

IT 915969-19-6DP, hydrogenated
(manufacture of metal hydride complexes useful as catalysts for hydrogenation of ring-opening polymerization polymers of cycloolefins with freedom from gel complication)

RN 915969-19-6 HCAPLUS

CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-

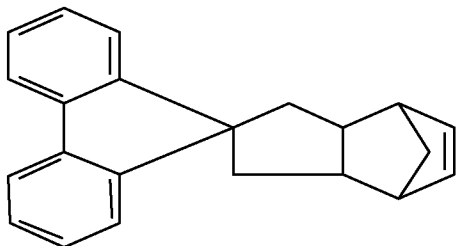
10/556,717

octahydro-2-methyl-, methyl ester, polymer with 1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] (CA INDEX NAME)

CM 1

CRN 908004-25-1

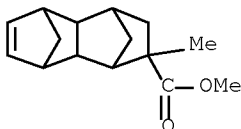
CMF C22 H20



CM 2

CRN 58732-15-3

CMF C15 H20 O2



CC 37-3 (Plastics Manufacture and Processing)

Section cross-reference(s): 67, 78

IT 123303-71-9DP, hydrogenated 133126-13-3DP, hydrogenated

666860-23-7DP, hydrogenated 915969-19-6DP, hydrogenated

(manufacture of metal hydride complexes useful as catalysts for hydrogenation of ring-opening polymerization polymers of cycloolefins with freedom from gel complication)

REFERENCE COUNT: 15 THERE ARE 15 CITED REFERENCES AVAILABLE FOR THIS RECORD. ALL CITATIONS AVAILABLE IN THE RE FORMAT

L37 ANSWER 10 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:1030832 HCAPLUS Full-text

DOCUMENT NUMBER: 145:397250

TITLE: Process for the preparation of spiro compound

INVENTOR(S): Kawashima, Naoyuki; Miyaki, Nobuyuki; Miyamoto, Yoshikazu; Kajiwara, Ichiro

PATENT ASSIGNEE(S): Jsr Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 22pp.

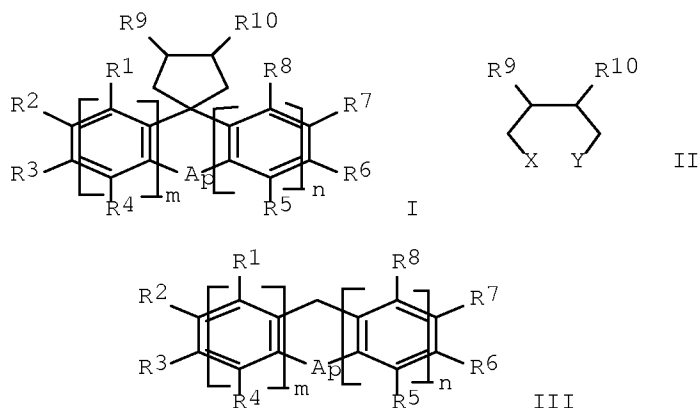
CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 1
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2006265176	A	20061005	JP 2005-85961	20050324
PRIORITY APPLN. INFO.:			JP 2005-85961	20050324

OTHER SOURCE(S): CASREACT 145:397250; MARPAT 145:397250
 ED Entered STN: 05 Oct 2006
 GI

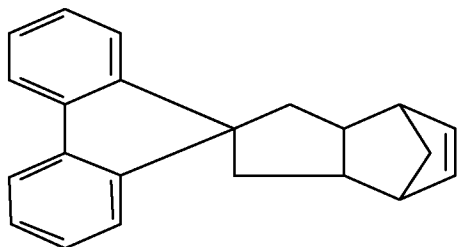


AB A process for the preparation of title compds. of formula I [R1-R8 = H or halo; R9R10 = (hetero)cyclyl; A = a single bond or divalent; m, n, p = 0-2] comprising cyclocondensation reaction of a compound of formula II (X, Y = halo or phosphate ester) with a compound of formula III [R1-R8 and A are defined as above] in the presence of metal hydride as bases is disclosed. The manufacturing method which can obtain the spiro compound cheaply and efficiently is provided.

IT 908004-25-1P
 (preparation of spiro compound by cyclocondensation of fluorene with 5,6-di(p-toluenesulfonyloxymethyl)bicyclo[2.2.1]hept-2-ene)

RN 908004-25-1 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro- (CA INDEX NAME)



CC 25-27 (Benzene, Its Derivatives, and Condensed Benzenoid Compounds)
 IT 908004-25-1P
 (preparation of spiro compound by cyclocondensation of fluorene with
 5,6-di(p-toluenesulfonyloxymethyl)bicyclo[2.2.1]hept-2-ene)

L37 ANSWER 11 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:886278 HCAPLUS Full-text

DOCUMENT NUMBER: 145:293498

TITLE: Preparation of norbornene derivatives by
 Diels-Alder reaction of cycloolefins and
 cyclopentadienes

INVENTOR(S): Kawashima, Naoyuki; Miyaki, Nobuyuki; Miyamoto,
 Yoshikazu; Kajiwara, Ichiro

PATENT ASSIGNEE(S): JSR Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 19pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

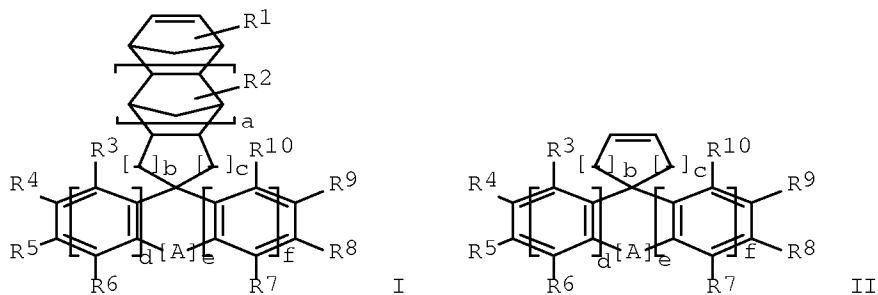
PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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JP 2006225316	A	20060831	JP 2005-40826	20050217
PRIORITY APPLN. INFO.:			JP 2005-40826	20050217

OTHER SOURCE(S): MARPAT 145:293498

ED Entered STN: 31 Aug 2006

GI



AB The derivs. I [R1-R10 = H, halo, C1-10 hydrocarbylene, etc.; a-f = 0-2 (b = c ≠ 0); A = single bond, O, S, SO, etc.], useful for index anisotropy-controlling monomers for optical-grade transparent resins, are prepared by Diels-Alder reaction of cycloolefins II (R1-R10, a-f = the same as above) with cyclopentadienes (at 100-250°). Thus, 0.12 mol cis-1,4-dichloro-2-butene was reacted with 0.1 mol fluorene in DMSO in the presence of NaOH and Et3BzNCl at 95-100° to give a product mixture, which was further reacted with dicyclopentadiene at 210° and recrystd. from MeOH to afford spiro[fluorene-9,8'-tricyclo[4.3.0.12,5][3]decene] in 22% yield.

IT 908004-25-1P

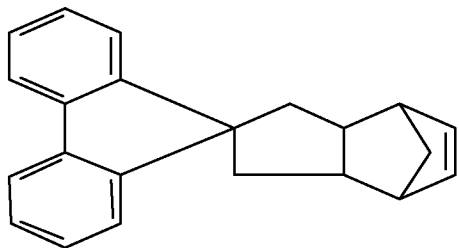
(monomers; preparation of norbornene monomers by Diels-Alder reaction of

10/556,717

cycloolefins and cyclopentadienes for optical-grade resin materials)

RN 908004-25-1 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro- (CA INDEX NAME)



CC 35-2 (Chemistry of Synthetic High Polymers)
Section cross-reference(s): 73

IT 908004-25-1P
(monomers; preparation of norbornene monomers by Diels-Alder reaction of cycloolefins and cyclopentadienes for optical-grade resin materials)

L37 ANSWER 12 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2006:705949 HCAPLUS Full-text

DOCUMENT NUMBER: 145:155751

TITLE: Optical films containing norbornene (co)polymers with positive wavelength dependency, manufacture of same films, and sheet polarizers comprising same films as protective films

INVENTOR(S): Miyaki, Nobuyuki; Miyamoto, Yoshikazu; Kawashima, Naoyuki; Yoshida, Shuichi; Kajiwara, Ichiro; Komiya, Zen

PATENT ASSIGNEE(S): Jsr Ltd., Japan

SOURCE: Jpn. Kokai Tokkyo Koho, 54 pp.

CODEN: JKXXAF

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 1

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
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JP 2006188671	A	20060720	JP 2005-351865	20051206
PRIORITY APPLN. INFO.:			JP 2004-355532	A 20041208

ED Entered STN: 20 Jul 2006

GI

* STRUCTURE DIAGRAM TOO LARGE FOR DISPLAY - AVAILABLE VIA OFFLINE PRINT *

AB Claimed are the optical films made of norbornene (co)polymers having structural unit I [m, n = 0-2; X = CH:CH, CH₂CH₂; R₁-8 = H, halo,

10/556,717

(substituted) C1-30 hydrocarbyl which may bear connecting group bearing (O, N, S, or Si), polar group; p, q, r = 0-2]. The optical films are manufactured by solvent casting or melt extrusion. The (co)polymers show stable pos. wavelength dependency in whole region at 400-800 nm regardless of changes of environmental temperature and moisture.

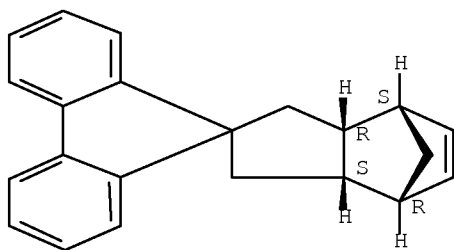
IT 791813-28-0F

(in preparation of norbornene monomers for metathetic ring-opening polymerization for manufacture of optical film)

RN 791813-28-0 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'S,7'R,7'aS)-rel- (CA INDEX NAME)

Relative stereochemistry.



IT 797785-12-7DP, hydrogenated

(metathetic ring-opening polymerized, optical film; manufacture of norbornene (co)polymer optical film, and sheet polarizer comprising same as protective film)

RN 797785-12-7 HCAPLUS

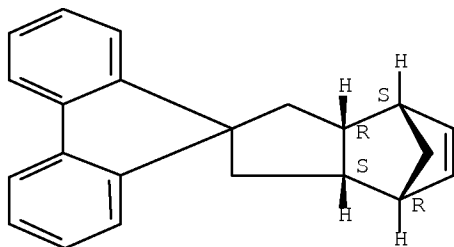
CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-octahydro-2-methyl-, methyl ester, polymer with rel-(3'aR,4'S,7'R,7'aS)-1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] (9CI) (CA INDEX NAME)

CM 1

CRN 791813-28-0

CMF C22 H20

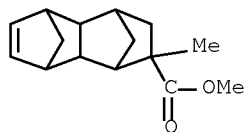
Relative stereochemistry.



CM 2

CRN 58732-15-3

CMF C15 H20 O2



CC 73-11 (Optical, Electron, and Mass Spectroscopy and Other Related Properties)
 Section cross-reference(s): 25, 38

IT 791813-28-0P
 (in preparation of norbornene monomers for metathetic ring-opening polymerization for manufacture of optical film)

IT 797785-12-7DP, hydrogenated
 (metathetic ring-opening polymerized, optical film; manufacture of norbornene (co)polymer optical film, and sheet polarizer comprising same as protective film)

L37 ANSWER 13 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN

ACCESSION NUMBER: 2004:1015980 HCAPLUS Full-text

DOCUMENT NUMBER: 142:7275

TITLE: Norbornene derivative, norbornene polymer produced by ring-opening (co)polymerization, and ring-opening (co)polymerization process

INVENTOR(S): Miyamoto, Yoshikazu; Miyaki, Nobuyuki; Goto, Kohei; Hashiguchi, Yuichi

PATENT ASSIGNEE(S): JSR Corporation, Japan

SOURCE: PCT Int. Appl., 83 pp.
 CODEN: PIXXD2

DOCUMENT TYPE: Patent

LANGUAGE: Japanese

FAMILY ACC. NUM. COUNT: 2

PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
WO 2004101478	A1	20041125	WO 2004-JP6095	20040427
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
JP 2004323489	A	20041118	JP 2003-135702	20030514
EP 1623967	A1	20060208	EP 2004-729707	20040427
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
JP 2005036201	A	20050210	JP 2004-172229	20040610
US 20070065747	A1	20070322	US 2005-556717	20051114

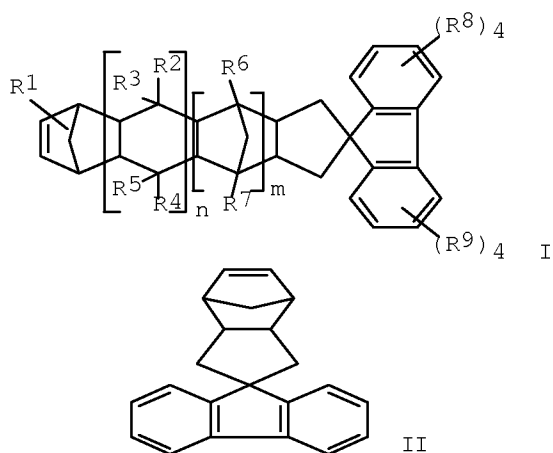
PRIORITY APPLN. INFO.:

JP 2003-135702	A	20030514
JP 2003-185203	A	20030627
JP 2003-60061	A	20030306
WO 2004-JP6095	W	20040427

OTHER SOURCE(S): MARPAT 142:7275

ED Entered STN: 25 Nov 2004

GI



AB Spiro norbornene derivative I [R1-R9 = H, halo, polar group, (un)substituted C1-30 alkyl which may have linking groups bearing O, N, S or Si atoms; s, t, u = 0-3; m, n = 0-2] is prepared and used as a precursor monomer for producing cycloolefin polymers having excellent transparency and heat resistance, reduced water absorption, and high solubility in organic solvents. Thus, exo II was prepared and polymerized with 8-methoxycarbonyl-8-methyltetracyclo[4.4.0.12,5.17,10]-3-dodecene to give a copolymer, which was hydrogenated and made into a transparent film showing peculiar birefringence and wavelength dependence.

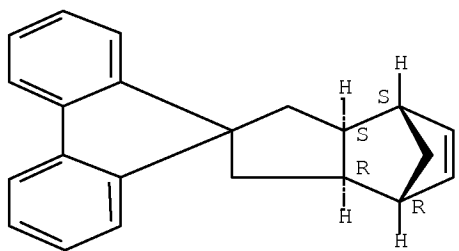
IT 791813-27-9P 791813-28-0P

(preparation and ring-opening polymerization of spiro norbornene derivs.)

RN 791813-27-9 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'R,7'S,7'aS)-rel- (CA INDEX NAME)

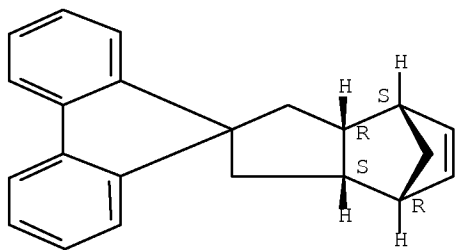
Relative stereochemistry.



RN 791813-28-0 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'S,7'R,7'aS)-rel- (CA INDEX NAME)

Relative stereochemistry.



IT 797785-11-6DP, hydrogenated 797785-12-7DP, hydrogenated

(preparation of spironorbornene polymers with good transparency and solubility and peculiar birefringence and wavelength dependence)

RN 797785-11-6 HCAPLUS

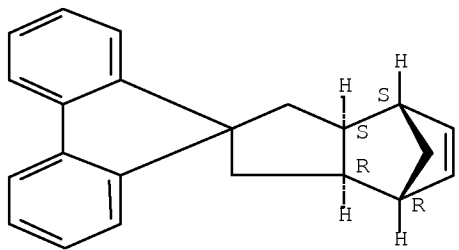
CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-octahydro-2-methyl-, methyl ester, polymer with rel-(3'aR,4'R,7'S,7'aS)-1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] (9CI) (CA INDEX NAME)

CM 1

CRN 791813-27-9

CMF C22 H20

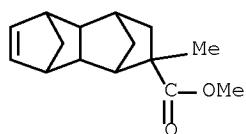
Relative stereochemistry.



CM 2

CRN 58732-15-3

CMF C15 H20 O2



RN 797785-12-7 HCAPLUS

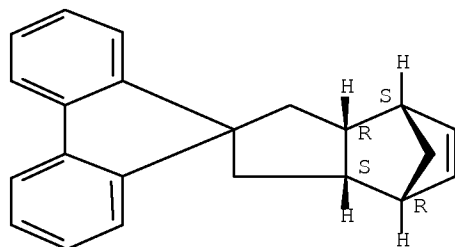
CN 1,4:5,8-Dimethanonaphthalene-2-carboxylic acid, 1,2,3,4,4a,5,8,8a-octahydro-2-methyl-, methyl ester, polymer with rel-(3'aR,4'S,7'R,7'aS)-1',3',3'a,4',7',7'a-hexahydrospiro[9H-fluorene-9,2'-[4,7]methano[2H]indene] (9CI) (CA INDEX NAME)

CM 1

CRN 791813-28-0

CMF C22 H20

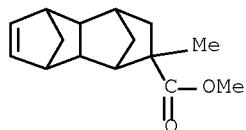
Relative stereochemistry.



CM 2

CRN 58732-15-3

CMF C15 H20 O2



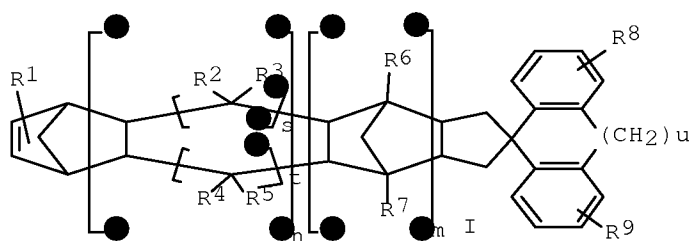
IC ICM C07C013-72

ICS C08G061-06
 CC 37-3 (Plastics Manufacture and Processing)
 Section cross-reference(s): 35
 IT 791813-27-9P 791813-28-0P
 (preparation and ring-opening polymerization of spiro norbornene derivs.)
 IT 797785-11-6DP, hydrogenated 797785-12-7DP,
 hydrogenated
 (preparation of spironorbornene polymers with good transparency and
 solubility and peculiar birefringence and wavelength dependence)
 REFERENCE COUNT: 3 THERE ARE 3 CITED REFERENCES AVAILABLE FOR
 THIS RECORD. ALL CITATIONS AVAILABLE IN THE
 RE FORMAT

L37 ANSWER 14 OF 14 HCAPLUS COPYRIGHT 2008 ACS on STN
 ACCESSION NUMBER: 2004:993154 HCAPLUS Full-text
 DOCUMENT NUMBER: 141:411390
 TITLE: Norbornenes having spiro structures for
 cycloolefin polymers showing desired optical
 properties
 INVENTOR(S): Miyamoto, Yoshikazu; Miyaki, Nobuyuki; Hashiguchi,
 Yuichi; Goto, Kohei
 PATENT ASSIGNEE(S): JSR Ltd., Japan
 SOURCE: Jpn. Kokai Tokkyo Koho, 29 pp.
 CODEN: JKXXAF
 DOCUMENT TYPE: Patent
 LANGUAGE: Japanese
 FAMILY ACC. NUM. COUNT: 2
 PATENT INFORMATION:

PATENT NO.	KIND	DATE	APPLICATION NO.	DATE
JP 2004323489	A	20041118	JP 2003-135702	20030514
WO 2004101478	A1	20041125	WO 2004-JP6095	20040427
W: AE, AG, AL, AM, AT, AU, AZ, BA, BB, BG, BR, BW, BY, BZ, CA, CH, CN, CO, CR, CU, CZ, DE, DK, DM, DZ, EC, EE, EG, ES, FI, GB, GD, GE, GH, GM, HR, HU, ID, IL, IN, IS, KE, KG, KP, KR, KZ, LC, LK, LR, LS, LT, LU, LV, MA, MD, MG, MK, MN, MW, MX, MZ, NA, NI, NO, NZ, OM, PG, PH, PL, PT, RO, RU, SC, SD, SE, SG, SK, SL, SY, TJ, TM, TN, TR, TT, TZ, UA, UG, US, UZ, VC, VN, YU, ZA, ZM, ZW RW: BW, GH, GM, KE, LS, MW, MZ, NA, SD, SL, SZ, TZ, UG, ZM, ZW, AM, AZ, BY, KG, KZ, MD, RU, TJ, TM, AT, BE, BG, CH, CY, CZ, DE, DK, EE, ES, FI, FR, GB, GR, HU, IE, IT, LU, MC, NL, PL, PT, RO, SE, SI, SK, TR, BF, BJ, CF, CG, CI, CM, GA, GN, GQ, GW, ML, MR, NE, SN, TD, TG				
EP 1623967	A1	20060208	EP 2004-729707	20040427
R: AT, BE, CH, DE, DK, ES, FR, GB, GR, IT, LI, LU, NL, SE, MC, PT, IE, SI, FI, RO, CY, TR, BG, CZ, EE, HU, PL, SK				
CN 1787983	A	20060614	CN 2004-80013152	20040427
US 20070065747	A1	20070322	US 2005-556717	20051114
PRIORITY APPLN. INFO.:			JP 2003-60061	A 20030306
			JP 2003-135702	A 20030514
			JP 2003-185203	A 20030627
			WO 2004-JP6095	W 20040427

ED Entered STN: 19 Nov 2004
 GI



AB The norbornenes are I [R1-R9 = H, halo, (O-, N-, S-, or Si-containing linkage-containing) C1-30 hydrocarbyl, polar group; s, t, u = 0-3; m, n = 0-2]. Thus, 5-norbornene-2-exo-3-exo-dimethanol was treated with p-MeC6H4SO2Cl in the presence of pyridine and treated with fluorene in the presence of BuLi to give exo-I (R1-R9 = H, m = n = u = 0).

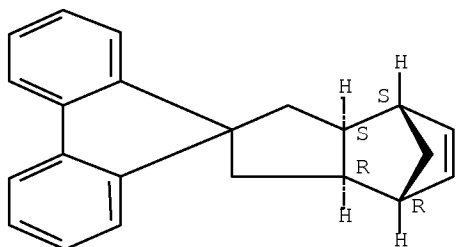
IT 791813-27-9P 791813-28-0P

(manufacture of norbornenes having spiro structures for cycloolefin polymers showing desired optical properties)

RN 791813-27-9 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'R,7'S,7'aS)-rel- (CA INDEX NAME)

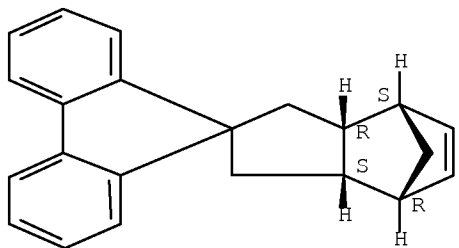
Relative stereochemistry.



RN 791813-28-0 HCAPLUS

CN Spiro[9H-fluorene-9,2'-[4,7]methano[2H]indene], 1',3',3'a,4',7',7'a-hexahydro-, (3'aR,4'S,7'R,7'aS)-rel- (CA INDEX NAME)

Relative stereochemistry.



IC ICM C07C013-72
ICS C08G061-08; G02B005-30; G02B006-00; G02B006-12; G02F001-1333;
G02F001-1335; G02F001-1336
CC 35-2 (Chemistry of Synthetic High Polymers)
Section cross-reference(s): 25
IT 791813-27-9P 791813-28-0P
(manufacture of norbornenes having spiro structures for cycloolefin
polymers showing desired optical properties)

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(FILE 'HOME' ENTERED AT 13:13:14 ON 14 JUL 2008)

FILE 'HCAPLUS' ENTERED AT 13:13:25 ON 14 JUL 2008

L1 2 SEA ABB=ON PLU=ON US20070065747/PN
SEL RN

FILE 'REGISTRY' ENTERED AT 13:13:38 ON 14 JUL 2008

L2 10 SEA ABB=ON PLU=ON (23067-44-9/BI OR 33493-88-8/BI OR
699-97-8/BI OR 791813-27-9/BI OR 791813-28-0/BI OR
86-73-7/BI OR 699-95-6/BI OR 797785-11-6/BI OR 797785-12-7/
BI OR 98-59-9/BI)
L3 STR
L4 0 SEA SSS SAM L3
L5 STR L3
L6 49 SEA SSS SAM L5
L7 STR L3
L8 5 SEA SSS SAM L7
L9 STR L7
L10 0 SEA SSS SAM L9
L11 3148 SEA SSS FUL L5
SAV L11 TEMP TRU717/A
L12 4 SEA ABB=ON PLU=ON L11 AND L2
L13 0 SEA SUB=L11 SSS SAM L9
L14 15 SEA SUB=L11 SSS FUL L9
SAV L14 TEMP TRU717A/A

FILE 'HCAPLUS' ENTERED AT 13:29:32 ON 14 JUL 2008

L15 14 SEA ABB=ON PLU=ON L14

FILE 'CAOLD' ENTERED AT 13:29:46 ON 14 JUL 2008

L16 0 SEA ABB=ON PLU=ON L14

FILE 'BEILSTEIN' ENTERED AT 13:29:55 ON 14 JUL 2008

L17 0 SEA ABB=ON PLU=ON L14
L18 0 SEA ABB=ON PLU=ON L14

FILE 'MARPAT' ENTERED AT 13:30:09 ON 14 JUL 2008

L19 0 SEA SSS SAM L9
L20 2 SEA SSS FUL L9

FILE 'REGISTRY' ENTERED AT 13:53:24 ON 14 JUL 2008

L21 STR L3
DIS SIA
L22 50 SEA SUB=L11 SSS SAM L21
L23 2 SEA SUB=L11 SSS SAM L3
L24 54 SEA SUB=L11 SSS FUL L3
L25 11 SEA ABB=ON PLU=ON L24 AND PMS/CI
L26 46 SEA ABB=ON PLU=ON L24 NOT FULLER?
L27 35 SEA ABB=ON PLU=ON L26 NOT FLUORANTHENE?
L28 34 SEA ABB=ON PLU=ON L27 NOT 1-100/SI
L29 30 SEA ABB=ON PLU=ON L28 NOT DISPIRO?
L30 28 SEA ABB=ON PLU=ON L29 NOT FURAN?
L31 4 SEA ABB=ON PLU=ON L30 NOT FLUORENE?
L32 29 SEA ABB=ON PLU=ON L29 NOT MORPHOLINYL?
L33 28 SEA ABB=ON PLU=ON L32 NOT MORPHOLINE?
L34 15 SEA ABB=ON PLU=ON L33 AND L14

10/556,717

L35 13 SEA ABB=ON PLU=ON L33 NOT L34

FILE 'HCAPLUS' ENTERED AT 14:12:08 ON 14 JUL 2008

L36 14 SEA ABB=ON PLU=ON L34

L37 14 SEA ABB=ON PLU=ON L15 OR L36

FILE 'MARPAT' ENTERED AT 14:12:30 ON 14 JUL 2008

L38 0 SEA ABB=ON PLU=ON L20 NOT L37